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SILYLATED TETRAHYDROFURAN DERIVATIVES

by

Curtis L. Schilling, Jr.

Prepared for Publication in  
Organometallics

Union Carbide Corporation  
Tarrytown, New York 10591

3 December 1982

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## SILYLATED TETRAHYDROFURAN DERIVATIVES

Curtis L. Schilling, Jr.  
Union Carbide Corporation  
Silicones 220, Technical Center  
Tarrytown, New York 10591

### ABSTRACT

2-Trimethylsilyltetrahydrofuran and 2-(2-trimethylsilylethyl)tetrahydrofuran have been isolated from dechlorinations of mixtures of silane monomers by potassium metal in tetrahydrofuran solvent. These low yield products may involve the common intermediacy of the 2-tetrahydrofuryl anion or the 2-tetrahydrofuryl radical and its trapping by trimethylchlorosilane and vinyltrimethylsilane.

### INTRODUCTION

Numerous reactions have been run in tetrahydrofuran (THF) solvent wherein products which arise by hydrogen abstraction from the solvent are isolated. For example, treatment of gem-dibromocyclopropanes with methylmagnesium bromide yields monobromocyclopropanes<sup>1</sup>. Grignard<sup>2,3</sup> or lithium<sup>4</sup> reactions of organopolyhalides with trimethylchlorosilane yield products in which halogens have been replaced by hydrogens. Triphenylsilyllithium<sup>5</sup> and  $\alpha,\omega$ -dilithio-perphenylpolysilanes<sup>6</sup> react with dichloromethane to yield triphenylmethylsilane and  $\alpha,\omega$ -dimethylperphenylpolysilanes, respectively.

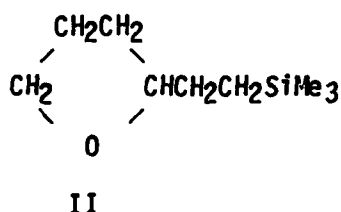
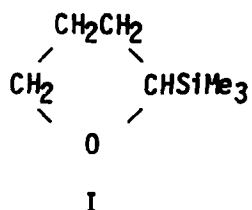
2-Lithiotetrahydrofuran is proposed as a product of the reaction of n-butyllithium with THF<sup>7,8</sup> with mass spectral evidence for monodeuterotetrahydrofuran when the reaction is quenched with D<sub>2</sub>O<sup>7</sup>. Deprotonation of THF by a bicyclononadienyl anion has also been reported<sup>9</sup>.

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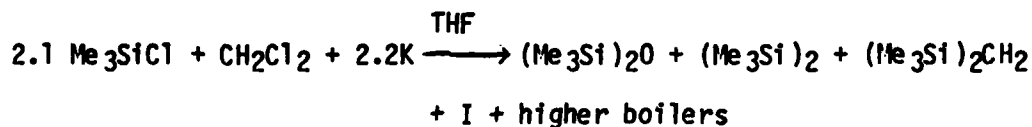
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Active metal dechlorinations of mixtures of silane monomers, including chlorosilanes, vinylic silanes, and chloromethyl silanes have yielded polycarbosilane precursors for silicon carbide.<sup>10</sup> Certain of these reactions, run using potassium metal in THF as the dechlorinating medium, have provided 2-trimethylsilyltetrahydrofuran (I) or 2-(2-trimethylsilyl)ethyl-tetrahydrofuran (II) in relatively low yields, in addition to the expected products.



Isolable quantities of I were obtained from the reaction of trimethylchlorosilane with dichloromethane using K metal in THF. Addition of a mixture of  $\text{Me}_3\text{SiCl}/\text{CH}_2\text{Cl}_2$  to refluxing THF/molten K metal suspension



at a rate maintaining the reflux temperature above 64°, followed by cooling, cautious termination with aqueous THF, and neutralization with conc. HCl yielded a group of products including I (5.3% yield based on  $\text{Me}_3\text{SiCl}$ ).

Preparative gas chromatography yielded a pure sample with appropriate NMR/mass spectra and elemental analyses ( $\text{C}_7\text{H}_{16}\text{OSi}$ , Calc'd: % C, 58.33, % H, 11.11, % Si, 19.44; Found: % C, 58.39, % H, 11.15, % Si, 19.66).

Similarly, dechlorination of ethyltrichlorosilane in the presence of vinyltrimethylsilane



yields a branched polycarbosilane + II (0.12% yield). A purified sample provided correct NMR/mass spectra and elemental analyses ( $\text{C}_9\text{H}_{20}\text{OSi}$ , Calc'd: % C, 62.79, % H, 11.63, % Si, 16.28; Found: % C, 63.03, % H, 11.87, % Si, 16.04. Yields of II up to 2.9% have been obtained from other reactions.

The isolations of I and II support the intermediacy of the 2-tetrahydrofuryl anion or the 2-tetrahydrofuryl radical. The anionic intermediate is more likely in the formation of I via simple displacement of chloride from  $\text{Me}_3\text{SiCl}$ . Compound II could form via a free radical process, or by Michael addition of an anionic intermediate to  $\text{CH}_2=\text{CHSiMe}_3$ . Both radical and anionic intermediates may be involved in a common pathway with K metal transforming 2-tetrahydrofuryl radicals to 2-tetrahydrofuryl anions by electron transfer. Our attempts to prepare I by quenching an *n*-butyllithium/THF solution<sup>7</sup> with  $\text{Me}_3\text{SiCl}$  were unsuccessful, while treatment of a refluxing solution of THF/ $\text{CH}_2=\text{CHSiMe}_3$  with benzoyl peroxide or azobisisobutyronitrile (method of reference 11) provided a very low yield of II.

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